

This listing of claims replaces all prior versions and listings of claims in this application.

**LISTING OF CLAIMS:**

Claim 1 (Currently Amended): A tablet prepared by direct compression of comprising crystals of a pharmaceutically acceptable salt of citalopram and pharmaceutically acceptable excipients, wherein the median particle size of the crystals is at least 40  $\mu\text{m}$ , which is prepared by direct compression of the pharmaceutically acceptable salt and pharmaceutically acceptable excipients.

Claims 2-3 (Canceled).

Claim 4 (Previously Presented): The tablet according to claim 1 which does not contain a binder.

Claim 5 (Previously Presented): The tablet according to claim 1 which contains 2-60% w/w active ingredient calculated as citalopram base.

Claim 6 (Previously Presented): The tablet according to claim 1 which contains a filler selected from lactose, sugars, calcium phosphates, starch, modified starches, microcrystalline cellulose, calcium sulfate and calcium carbonate.

**Claim 7 (Previously Presented):** The tablet according to claim 6, wherein the filler is a microcrystalline cellulose.

**Claim 8 (Previously Presented):** The tablet according to claim 1 which contains a lubricant selected from metallic stearates, stearic acid, wax, hydrogenated vegetable oil, talc and colloidal silica.

**Claim 9 (Previously Presented):** The tablet according to claim 8, wherein the lubricant is magnesium stearate or calcium stearate.

**Claim 10 (Previously Presented):** The tablet according to claim 1 which is substantially free of lactose.

**Claim 12 (Previously Presented):** The tablet according to claim 1 wherein the pharmaceutically acceptable salt is citalopram hydrobromide or citalopram hydrochloride.

**Claim 13 (Previously Presented):** The tablet according to claim 12, wherein the pharmaceutically acceptable salt is citalopram hydrobromide.

**Claims 14-35 (Canceled).**

**Claim 36 (Previously Presented):** The tablet of claim 1, which contains 10-40% w/w active ingredient calculated as citalopram base.

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**Claim 37 (Previously Presented):** The tablet of claim 1, which contains 15-25% w/w active ingredient calculated as citalopram base.

**Claim 38 (Previously Presented):** The tablet of claim 6, wherein said filler is a sugar selected from the group consisting of sorbitol, mannitol, dextrose and sucrose.

**Claim 39 (Previously Presented):** The tablet of claim 6, wherein said filler is a calcium phosphate selected from the group consisting of dibasic, tribasic, hydrous and anhydrous calcium phosphate.

**Claim 40 (Previously Presented):** The tablet of claim 8, wherein said lubricant is a metallic stearate selected from the group consisting of magnesium, calcium and sodium stearate.

**Claim 41 (Previously Presented):** The tablet of claim 1, wherein the crystals have a median particle size of 40-200  $\mu\text{m}$ .

**Claim 42 (Previously Presented):** The tablet of claim 1, wherein the crystals have a median particle size of 45-150  $\mu\text{m}$ .

**Claim 43 (Previously Presented):** The tablet of claim 1, wherein the crystals have a median particle size of 50-100  $\mu\text{m}$ .

Claims 44-92 (Canceled).

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